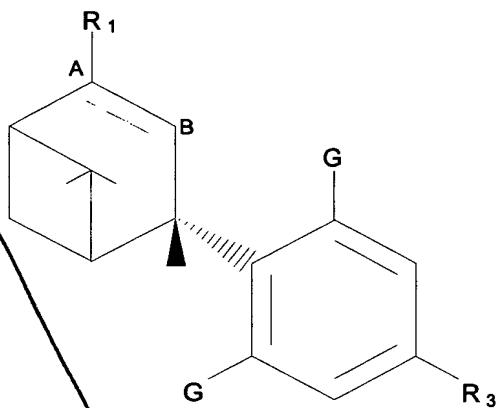


THE CLAIMS

What is claimed is:

- 5 1. A pharmaceutical composition for treating or preventing hypertension, inflammation, peripheral pain, gastrointestinal disorders, or autoimmune diseases, comprising as an active ingredient a compound of the general formula:



- having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer,
10 wherein:
the dashed line A---B designates an optional double bond,
15 R_1 is (a) $-R'N(R'')_2$ wherein R' is C_1-C_5 straight or branched chain alkyl and each R'' , which may be the same or different, is hydrogen or C_1-C_5 straight or branched chain alkyl optionally containing a terminal $-OR'''$ or $-OC(O)R'''$ moiety wherein R''' is hydrogen or C_1-C_5 straight or branched chain alkyl, (b) $-Q$ wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) $-R'X$ wherein R' is C_1-C_5 straight or branched chain alkyl and X is halogen, (d) $-R'C(O)N(R'')_2$ wherein R' is a direct bond or C_1-C_5 straight or branched chain alkyl and each R'' , which may be the same or different, is hydrogen or C_1-C_5 straight or branched chain alkyl optionally containing a terminal $-OR'''$ or $-OC(O)R'''$ moiety wherein R''' is hydrogen or C_1-C_5 straight or branched chain alkyl, (e) $-R'C(O)OR''$ wherein R' is a direct bond or C_1-C_5 straight or branched chain alkyl and R'' is hydrogen or C_1-C_5 straight or branched chain alkyl optionally containing a terminal $-OR'''$ or
20

~~-OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R'~~
~~wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight~~
~~or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;~~

~~G is hydrogen; and~~

~~5 R₃ is (a) C₁-C₁₂ straight or branched chain alkyl, (b) -OR''', in which R''' is a straight
chain or branched C₂-C₉ alkyl which may be substituted at the terminal carbon atom by a phenyl
group, or (c) -(CH₂)_nOR''' wherein n is an integer of 1 to 7 and R''' is hydrogen or
C₁-C₅ alkyl.~~

~~10 2. The compound of claim 1, wherein R₃ is a straight or branched chain C₅-C₁₂ alkyl.~~

~~3. The compound of claim 1, wherein R₃ is 1,1-dimethyl heptyl or 1,2-dimethyl heptyl.~~

~~15 4. The compound of claim 1 wherein Q is a saturated or unsaturated ring of 4 to 8 members
consisting of C with at least one of N, S, and O, said ring being optionally substituted with -
COR''' or -COOR''' wherein R''' is a hydrogen or C₁-C₅ straight or branched chain alkyl.~~

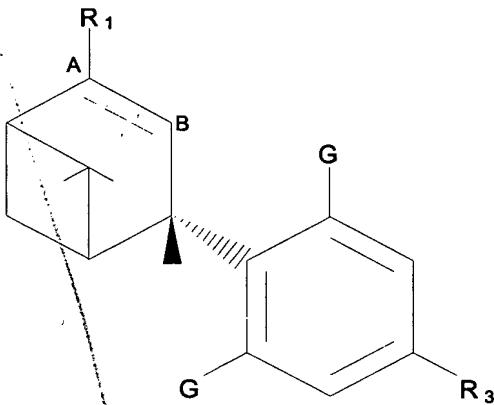
~~20 5. The compound of claim 1, wherein R₁ is -CH₂OH, -C(O)N(R'')₂, -C(O)OR'',
-COOH, an amino acid, or a carboxamide.~~

~~6. A pharmaceutical composition for treating, preventing, or managing hypertension,
inflammation, peripheral pain, gastrointestinal disorders, or autoimmune diseases comprising
as an active ingredient a therapeutically effective amount of a compound of claim 1.~~

~~25 7. The pharmaceutical composition of claim 6 further comprising a pharmaceutically
acceptable diluent or carrier.~~

~~8. The pharmaceutical composition of claim 7, wherein the diluent is an aqueous cosolvent
solution comprising a pharmaceutically acceptable cosolvent, a micellar solution or emulsion
30 prepared with natural or synthetic ionic or non-ionic surfactants, or a combination of such
cosolvent and micellar or emulsion solutions.~~

9. A method for preventing, treating, or managing hypertension, inflammation, peripheral pain, gastrointestinal disorders, or autoimmune diseases comprising administering to an individual in need thereof a pharmaceutical composition comprising a therapeutically effective amount a compound of the general formula:



- 5 having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer, wherein:
- A---B designates an optional double bond,
- R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl
- 10 optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different,
- 15 is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R'
- 20 wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;

G is hydrogen, halogen, or -OR₂ wherein R₂ is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR'', -OC(O)R'', C(O)OR'', or -C(O)R'' moiety wherein R'' is hydrogen or C₁-C₅ straight or branched chain alkyl; and

R₃ is (a) C₁-C₁₂ straight or branched chain alkyl, (b) -OR''', in which R''' is a straight 5 chain or branched C₂-C₉ alkyl which may be substituted at the terminal carbon atom by a phenyl group, or (c) -(CH₂)_nOR''' wherein n is an integer of 1 to 7 and R''' is hydrogen or C₁-C₅ alkyl.

10. The method of claim 9 wherein, R₁ is -CH₂OH, G is hydrogen or OR₂, R₂ is a lower alkyl group, and R₃ is a straight or branched chain C₅-C₁₂ alkyl.

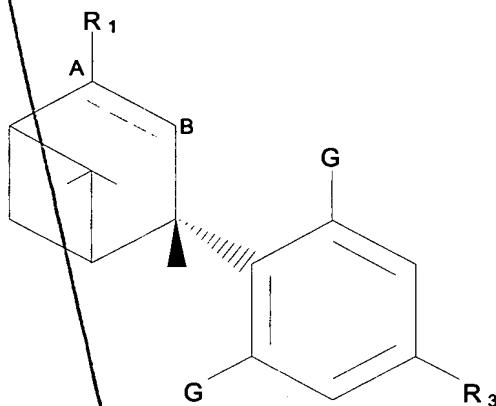
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11. The method of claim 10, wherein G is -OCH₃ and R₃ is 1,1-dimethyl heptyl.

12. The method of claim 10, wherein R₁ is -CH₂OH, G is -OCH₃, and R₃ is 1,1-dimethyl heptyl.

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13. A method for preventing, treating, or managing tumors expressing CB2 receptors comprising administering to an individual in need thereof a pharmaceutical composition comprising a therapeutically effective amount a compound of the general formula:



having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer,

20 wherein:

A---B designates an optional double bond,

R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R' wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;

15 G is hydrogen, halogen, or -OR₂ wherein R₂ is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR'', -OC(O)R'', C(O)OR'', or -C(O)R'' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl; and

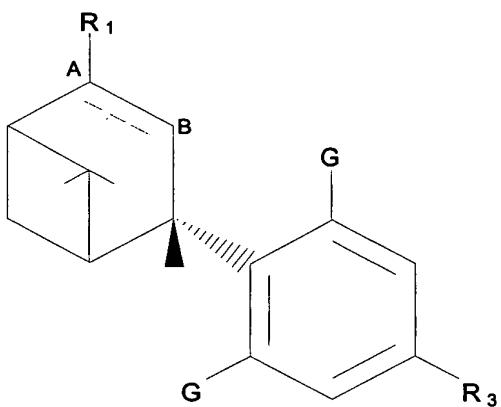
16 R₃ is (a) C₁-C₁₂ straight or branched chain alkyl, (b) -OR''', in which R''' is a straight chain or branched C₂-C₉ alkyl which may be substituted at the terminal carbon atom by a phenyl group, or (c) -(CH₂)_nOR''' wherein n is an integer of 1 to 7 and R''' is hydrogen or C₁-C₅ alkyl.

14. The method of claim 13 wherein, R₁ is -CH₂OH, G is hydrogen or OR₂, R₂ is a lower alkyl group, and R₃ is a straight or branched chain C₅-C₁₂ alkyl.

25 15. The method of claim 14, wherein G is -OCH₃ and R₃ is 1,1-dimethyl heptyl.

16. The method of claim 14, wherein R₁ is -CH₂OH, G is -OCH₃, and R₃ is 1,1-dimethyl heptyl.

30 17. A CB2 specific antagonist comprising a compound of the general formula:



having the (3S,4S) configuration, and which is essentially free of the (3R,4R) enantiomer, wherein:

A---B designates an optional double bond,

R₁ is (a) -R'N(R'')₂ wherein R' is C₁-C₅ straight or branched chain alkyl and each R'',

- 5 which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (b) -Q wherein Q is a heterocyclic moiety having a labile hydrogen atom so that said moiety acts as a carboxylic acid analogue, (c) -R'X wherein R' is C₁-C₅ straight or branched chain alkyl and X is halogen, (d) -R'C(O)N(R'')₂ wherein R' is a direct
- 10 bond or C₁-C₅ straight or branched chain alkyl and each R'', which may be the same or different, is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (e) -R'C(O)OR'' wherein R' is a direct bond or C₁-C₅ straight or branched chain alkyl and R'' is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''' or
- 15 -OC(O)R''' moiety wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl, (f) -R' wherein R' is C₁-C₅ straight or branched chain alkyl, or (g) -R'OR''' wherein R' is C₁-C₅ straight or branched chain alkyl and R''' is hydrogen or C₁-C₅ alkyl;

G is hydrogen, halogen, or -OR₂ wherein R₂ is hydrogen or C₁-C₅ straight or branched chain alkyl optionally containing a terminal -OR''', -OC(O)R''', C(O)OR''', or -C(O)R'' moiety

- 20 wherein R''' is hydrogen or C₁-C₅ straight or branched chain alkyl; and

~~R₃ is (a) C₁-C₁₂ straight or branched chain alkyl, (b) -OR'''', in which R''' is a straight chain or branched C₂-C₉ alkyl which may be substituted at the terminal carbon atom by a phenyl group, or (c) -(CH₂)_nOR''' wherein n is an integer of 1 to 7 and R''' is hydrogen or C₁-C₅ alkyl.~~

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